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CLAIMS

1. A method of treating a person with rheumatoid arthritis, said method comprising administering to said person an effective amount of a compound that blocks an interaction between DC-SIGN and ICAM-3.

- 2. The method of claim 1 wherein said compound is selected from the group consisting of a mannose carbohydrate, a fucose carbohydrate, a plant lectin, an antibiotic, a sugar, a protein and an antibody.
- 3. The method of claim 2 wherein said mannose carbohydrate is selected from the group consisting of mannan and D-mannose.
- 4. The method of claim 2 wherein said fucose carbohydrate is L-fucose.
- 5. The method of claim 2 wherein said plant lectin is concanavalin A.
- 6. The method of claim 2 wherein said antibiotic is pradimicin A.
- 7. The method of claim 2 wherein said sugar is selected from the group consisting of N-acetyl-D-glucosamine and galactose.
- 8. The method of claim 2 wherein said protein is selected from the group consisting of the HIV envelope glycoprotein gp120, an analog of gp120 that binds DC-SIGN, or a fragment of gp120 that binds DC-SIGN.
- 9. The method of claim 2 wherein said antibody is selected from the group consisting of a) an antibody or antibody fragment that binds to DC-SIGN and b) an antibody or antibody fragment that binds to ICAM-3.

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10. The method of claim 9 wherein said antibody is a humanized antibody, a chimeric antibody, a polyclonal antibody, a monoclonal antibody, or a single chain antibody.

- 11. The method of claim 9 wherein said antibody fragment is an Fab, an F(ab')₂, an F(ab'), an Fv or an Fd.
- 12. The method of claim 9 wherein said antibody is selected from the group consisting of AZN-D1 and AZN-D2.
- 13. The method of claim 2 wherein said compound is soluble DC-SIGN, a soluble analog of DC-SIGN that binds ICAM-3, or a soluble fragment of DC-SIGN that binds ICAM-3.
- 14. A method to diagnose whether a human has rheumatoid arthritis, said method comprising measuring the percentage of inflammatory cells that express DC-SIGN from synovia of said human, wherein an amount of 50% or greater of said cells expressing DC-SIGN is an indication of rheumatoid arthritis.
- 15. The method of claim 14 wherein said amount is 60% or greater of said cells.
- 16. The method of claim 14 wherein said amount is 75% or greater of said cells.
- 17. The method of claim 14 wherein said measuring is performed using an antibody that binds DC-SIGN.
- 18. The method of claim 17 wherein said antibody is AZN-D1 or AZN-D2.
- 19. A method to isolate macrophages from other cells in a biological sample, said method comprising a) contacting said sample with an agent that binds DC-SIGN and b) separating cells that bind to said agent from cells that do not bind to said agent.

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- 20. The method of claim 19 wherein said agent is an antibody.
- 21. The method of claim 20 wherein said antibody is AZN-D1 or AZN-D2.
- 22. The method of claim 19 wherein said agent is selected from the group consisting of a mannose carbohydrate, a fucose carbohydrate, a plant lectin, an antibiotic, a sugar, and a protein.
- 23. The method of claim 19 wherein said sample is obtained from person with rheumatoid arthritis.
- 24. The method according to claim 19 wherein said antibody is attached to a solid support.
- 25. The method according to claim 19 wherein said macrophages express CD68.